

This Page Is Inserted by IFW Operations  
and is not a part of the Official Record

## **BEST AVAILABLE IMAGES**

Defective images within this document are accurate representations of the original documents submitted by the applicant.

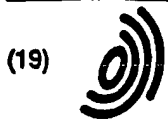
Defects in the images may include (but are not limited to):

- BLACK BORDERS
- TEXT CUT OFF AT TOP, BOTTOM OR SIDES
- FADED TEXT
- ILLEGIBLE TEXT
- SKEWED/SLANTED IMAGES
- COLORED PHOTOS
- BLACK OR VERY BLACK AND WHITE DARK PHOTOS
- GRAY SCALE DOCUMENTS

**IMAGES ARE BEST AVAILABLE COPY.**

**As rescanning documents *will not* correct images,  
please do not report the images to the  
Image Problems Mailbox.**

**THIS PAGE BLANK (USPTO)**



Europäisches Patentamt  
European Patent Office  
Office européen des brevets



F15

(11) EP 0 540 290 B1

(12) EUROPEAN PATENT SPECIFICATION

(45) Date of publication and mention  
of the grant of the patent:  
28.01.1998 Bulletin 1998/05

(51) Int. Cl.<sup>6</sup>: A61F 2/06, C23F 1/02

(21) Application number: 92309822.2

(22) Date of filing: 27.10.1992

(54) Expandable stents

Aufweitbare Stents

Stents expansibles

(84) Designated Contracting States:  
BE CH DE FR GB IT LI NL

(30) Priority: 28.10.1991 US 783558

(43) Date of publication of application:  
05.05.1993 Bulletin 1993/18

(60) Divisional application:  
97202436.8 / 0 807 424  
96201402.3 / 0 734 699

(73) Proprietor:  
ADVANCED CARDIOVASCULAR SYSTEMS, INC.  
Santa Clara California 95052-8167 (US)

(72) Inventors:  
• Lau, Lilip  
Cupertino, CA 95014 (US)  
• Hartigan, William M.  
Fremont, CA 94555 (US)

• Frantzen, John J.  
San Jose, CA 95132 (US)

(74) Representative:  
Mayes, Stuart David et al  
BOULT WADE TENNANT,  
27 Fumival Street  
London EC4A 1PQ (GB)

(56) References cited:

EP-A- 0 062 300	EP-A- 0 338 816
EP-A- 0 361 192	EP-A- 0 364 787
EP-A- 0 407 951	EP-A- 0 423 916
WO-A-91/07139	WO-A-92/09246
US-A- 4 733 665	US-A- 4 762 128
US-A- 4 767 418	US-A- 4 856 516
US-A- 4 994 071	US-A- 5 135 536

Remarks:

The file contains technical information submitted  
after the application was filed and not included in  
this specification

Note: Within nine months from the publication of the mention of the grant of the European patent, any person may give notice to the European Patent Office of opposition to the European patent granted. Notice of opposition shall be filed in a written reasoned statement. It shall not be deemed to have been filed until the opposition fee has been paid. (Art. 99(1) European Patent Convention).

## Description

This invention relates to expandable endoprosthesis devices, generally called stents, which are adapted to be implanted into a patient's body lumen, such as blood vessel, to maintain the patency thereof. These devices are very useful in the treatment of atherosclerotic stenosis in blood vessels.

Stents are generally tubular shaped devices which function to hold open a segment of a blood vessel or other anatomical lumen. They are particularly suitable for use to support and hold back a dissected arterial lining which can occlude the fluid passageway there-through.

Further details of prior art stents can be found in U.S. Patent 3,868,956 (Affidi *et al.*); U.S. Patent 4,512,338 (Balko *et al.*); U.S. Patent 4,553,545 (Maass *et al.*); U.S. Patent 4,733,665 (Palmaz); U.S. Patent 4,762,128 (Rosenbluth); U.S. Patent 4,800,882 (Gianturco); U.S. Patent 4,856,516 (Hillstead); and U.S. Patent 4,886,062 (Wiktor).

Various means have been described to deliver and implant stents. One method frequently described for delivering a stent to a desired intraluminal location includes mounting the expandable stent on an expandable member, such as a balloon, provided on the distal end of an intravascular catheter, advancing the catheter to the desired location within the patient's body lumen, inflating the balloon on the catheter to expand the stent into a permanent expanded condition and then deflating the balloon and removing the catheter. One of the difficulties encountered using prior stents involved maintaining the radial rigidity needed to hold open a body lumen while at the same time maintaining the longitudinal flexibility of the stent to facilitate its delivery.

What has been needed and heretofore unavailable is a stent which has a high degree of flexibility so that it can be advanced through tortuous passageways and can be readily expanded and yet have the mechanical strength to hold open the body lumen into which it is expanded.

One prior art device which goes somewhat towards addressing this problem is described in US-A-4,994,071. The described stent includes a trunk portion constructed from a series of generally parallel oriented loops interconnected by a series of half-hitch connections which extend along an axial dimension. In combination the series of loops form a generally cylindrical lattice or matrix.

The present invention is directed to an expandable stent which is relatively flexible along its longitudinal axis to facilitate delivery through tortuous body lumens, but which is stiff and stable enough radially in an expanded condition to maintain the patency of a body lumen such as an artery when implanted therein.

According to a first aspect of the present invention there is provided a longitudinally flexible stent formed from an elongated tubular member and comprising a

plurality of cylindrically shaped elements which are independently expandable in the radial direction and which are connected to one another by interconnecting elements so as to be generally aligned on a common longitudinal axis such that, other than at an end of the stent, each cylindrically shaped element has two adjacent cylindrically shaped elements spaced in opposite axial directions, the or each interconnecting element connecting one of said cylindrically shaped elements to one of said adjacent cylindrically shaped elements being circumferentially displaced from the or each interconnecting element connecting said cylindrically shaped element to the other of said adjacent cylindrically shaped elements.

Thus the stent of the invention includes a plurality of radially expandable cylindrical elements which are relatively independent in their ability to expand and to flex relative to one another. The individual radially expandable cylindrical elements of the stent are preferably dimensioned so as to be longitudinally shorter than their own diameters. Interconnecting elements or struts extend between adjacent cylindrical elements and provide increased stability and are preferably positioned to prevent warping of the stent upon the expansion thereof. The resulting stent structure is a series of radially expandable cylindrical elements which are spaced longitudinally close enough so that small dissections in the wall of a body lumen may be pressed back into position against the luminal wall, but not so close as to compromise the longitudinal flexibility of the stent. The individual cylindrical elements may rotate slightly relative to adjacent cylindrical elements without significant deformation, cumulatively giving a stent which is flexible along its length and about its longitudinal axis but which is still very stiff in the radial direction in order to resist collapse.

A stent embodying features of the invention can be readily delivered to the desired luminal location by mounting it on an expandable member of a delivery catheter, for example a balloon, and passing the catheter-stent assembly through the body lumen to the implantation site. A variety of means for securing the stent to the expandable member on the catheter for delivery to the desired location are available. It is presently preferred to compress the stent onto the balloon. Other means to secure the stent to the balloon include providing ridges or collars on the inflatable member to restrain lateral movement, or using bioresorbable temporary adhesives.

The presently preferred structure for the expandable cylindrical elements which form the stents of the present invention is to have a generally circumferential undulating pattern, e.g. serpentine. The transverse cross-section of the undulating component of the cylindrical element is relatively small and preferably has an aspect ratio of about two to one to about 0.5 to one. A one to one aspect ratio has been found particularly suitable. The open reticulated structure of the stent allows

for the perfusion of blood over a large portion of the arterial wall which can improve the healing and repair of a damaged arterial lining.

The radial expansion of the expandable cylinder deforms the undulating pattern thereof similar to changes in a waveform which result from decreasing the waveform's amplitude and the frequency. Preferably, the undulating patterns of the individual cylindrical structures are in phase with each other in order to prevent the contraction of the stent along its length when it is expanded. The cylindrical structures of the stent are plastically deformed when expanded (except with NiTi alloys) so that the stent will remain in the expanded condition and, therefore, they must be sufficiently rigid when expanded to prevent the collapse thereof in use. With superelastic NiTi alloys, the expansion occurs when the stress of compression is removed so as to allow the phase transformation from austenite back to martensite and as a result the expansion of the stent.

The elongated elements which interconnect adjacent cylindrical elements should have a transverse cross-section similar to the transverse dimensions of the undulating components of the expandable cylindrical elements. The interconnecting elements are formed in a unitary structure with the expandable cylindrical elements from the same intermediate product, namely a tubular element.

Preferably, all of the interconnecting elements of a stent are joined at either the peaks or the valleys of the undulating structure of the cylindrical elements which form the stent. In this manner there is no shortening of the stent upon expansion.

The number and location of elements interconnecting adjacent cylindrical elements can be varied in order to develop the desired longitudinal flexibility in the stent structure both in the unexpanded as well as the expanded condition. These properties are important to minimize alteration of the natural physiology of the body lumen into which the stent is implanted and to maintain the compliance of the body lumen which is internally supported by the stent. Generally, the greater the longitudinal flexibility of the stent, the easier and the more safely it can be delivered to the implantation site.

In a presently preferred arrangement

the stent is conveniently and easily formed by coating stainless steel hypotubing with a material resistant to chemical etching, and then removing portions of the coating to expose portions of underlying tubing which are to be removed to develop the desired stent structure. The exposed portions of the tubing are removed by chemically etching from the tubing exterior leaving the coated portion of the tubing material in the desired pattern of the stent structure. The etching process develops smooth openings in the tubing wall without burrs or other artifacts which are characteristic of mechanical or laser machining processes in the small sized products contemplated. Moreover, a computer controlled laser patterning process to remove the chem-

ical resistive coating makes photolithography technology adaptable to the manufacture of these small products. The forming of a mask in the extremely small sizes needed to make the small stents of the invention would be a most difficult task. A plurality of stents can be formed from one length of hypotubing by repeating the stent pattern and providing small webs or tabs to interconnect the stents. After the etching process, the stents can be separated by severing the small webs or tabs which connect them.

Other features and advantages of the present invention will become more apparent from the following detailed description of the invention, when taken in conjunction with the accompanying exemplary drawings.

FIG. 1 is an elevational view, partially in section, of a stent embodying features of the invention which is mounted on a delivery catheter and disposed within a damaged artery.

FIG. 2 is an elevational view, partially in section, similar to that shown in FIG. 1 wherein the stent is expanded within a damaged artery, pressing the damaged lining against the arterial wall.

FIG. 3 is an elevational view, partially in section showing the expanded stent within the artery after withdrawal of the delivery catheter.

FIG. 4 is a perspective view of a stent embodying features of the invention in an unexpanded state, with one end of the stent being shown in an exploded view to illustrate the details thereof.

FIG. 5 is a plan view of a flattened section of a stent of the invention which illustrates the undulating pattern of the stent shown in FIG. 4.

FIG. 6 is a schematic representation of equipment for selectively removing coating applied to tubing in the manufacturing of the stents of the present invention.

FIGS. 7 to 9 are perspective views schematically illustrating various configurations of interconnective element placement between the radially expandable cylindrical elements of the stent.

FIG. 10 is a plan view of a flattened section of a stent illustrating an alternate undulating pattern in the expandable cylindrical elements of the stent which are out of phase.

FIG. 1 illustrates a stent 10 incorporating features of the invention which is mounted onto a delivery catheter 11. The stent generally comprises a plurality of radially expandable cylindrical elements 12 disposed generally coaxially and interconnected by elements 13 disposed between adjacent cylindrical elements. The delivery catheter 11 has an expandable portion or balloon 14 for expanding the stent 10 within an artery 15. The artery 15, as shown in FIG. 1, has a dissected lining 16 which has occluded a portion of the arterial passageway.

The delivery catheter 11 onto which the stent 10 is mounted, is essentially the same as a conventional balloon dilatation catheter used for angioplasty procedures. The balloon 14 may be formed of suitable

materials such as polyethylene, polyethylene terephthalate, polyvinyl chloride, nylon and ionomers such as Surlyn™ manufactured by the Polymer Products Division of the Du Pont Company. Other polymers may also be used. In order for the stent 10 to remain in place on the balloon 14 during delivery to the site of the damage within the artery 15, the stent 10 is compressed onto the balloon. A retractable protective delivery sleeve 20 as described in US-A-5,507,768 and entitled "Stent Delivery System for use with Body Lumens in treating Endo-Arterial Prosthesis" may be provided to further ensure that the stent stays in place on the expandable portion of the delivery catheter 11 and prevent abrasion of the body lumen by the open surface of the stent 10 during delivery to the desired arterial location. Other means for securing the stent 10 onto the balloon 14 may also be used, such as providing collars or ridges on the ends of the working portion, *i.e.* the cylindrical portion, of the balloon.

Each radially expandable cylindrical element 12 of the stent 10 may be independently expanded. Therefore, the balloon 14 may be provided with an inflated shape other than cylindrical, *e.g.* tapered, to facilitate implantation of the stent 10 in a variety of body lumen shapes.

In a preferred embodiment, the delivery of the stent 10 is accomplished in the following manner. The stent 10 is first mounted onto the inflatable balloon 14 on the distal extremity of the delivery catheter 11. The balloon 14 is slightly inflated to secure the stent 10 onto the exterior of the balloon. The catheter-stent assembly is introduced within the patient's vasculature in a conventional Seldinger technique through a guiding catheter (not shown). A guidewire 18 is disposed across the damaged arterial section with the detached or dissected lining 16 and then the catheter-stent assembly is advanced over a guidewire 18 within the artery 15 until the stent 10 is directly under the detached lining 16. The balloon 14 of the catheter is expanded, expanding the stent 10 against the artery 15, which is illustrated in FIG. 2. While not shown in the drawing, the artery 15 is preferably expanded slightly by the expansion of the stent 10 to seat or otherwise fix the stent 10 to prevent movement. In some circumstances during the treatment of stenotic portions of an artery, the artery may have to be expanded considerably in order to facilitate passage of blood or other fluid therethrough.

The stent 10 serves to hold open the artery 15 after the catheter 11 is withdrawn, as illustrated by FIG. 3. Due to the formation of the stent 10 from an elongated tubular member, the undulating component of the cylindrical elements of the stent 10 is relatively flat in transverse cross-section, so that when the stent is expanded, the cylindrical elements are pressed into the wall of the artery 15 and as a result do not interfere with the blood flow through the artery 15. The cylindrical elements 12 of stent 10 which are pressed into the wall of the artery 15 will eventually be covered with endothelial cell

growth which further minimizes blood flow interference. The undulating portion of the cylindrical sections 12 provide good tacking characteristics to prevent stent movement within the artery. Furthermore, the closely spaced cylindrical elements 12 at regular intervals provide uniform support for the wall of the artery 15, and consequently are well adapted to tack up and hold in place small flaps or dissections in the wall of the artery 15 as illustrated in FIGS. 2 and 3.

FIG. 4 is an enlarged perspective view of the stent 10 shown in FIG. 1 with one end of the stent shown in an exploded view to illustrate in greater detail placement of interconnecting elements 13 between adjacent radially expandable cylindrical elements 12. Each pair of the interconnecting elements 13 on one side of a cylindrical element 12 are preferably placed to achieve maximum flexibility for a stent. In the embodiment shown in FIG. 4 the stent 10 has three interconnecting elements 13 between adjacent radially expandable cylindrical elements 12 which are 120 degrees apart. Each pair of interconnecting elements 13 on one end of a cylindrical element 12 are offset radially 60 degrees from the pair on the other side of the cylindrical element. The alternation of the interconnecting elements results in a stent which is longitudinally flexible in essentially all directions. Various configurations for the placement of interconnecting elements are possible, and several examples are illustrated schematically in FIGS. 7-9. However, as previously mentioned, all of the interconnecting elements of an individual stent should be secured to either the peaks or valleys of the undulating structural elements in order to prevent shortening of the stent during the expansion thereof.

FIG. 9 illustrates a stent of the present invention wherein three interconnecting elements 13 are disposed between radially expandable cylindrical elements 12. The interconnecting elements 13 are distributed radially around the circumference of the stent at a 120 degree spacing. Disposing four or more interconnecting elements between adjacent cylindrical elements 12 will generally give rise to the same considerations discussed above for two and three interconnecting elements.

The properties of the stent 10 may also be varied by alteration of the undulating pattern of the cylindrical elements 12. FIG. 10 illustrates an alternative stent structure in which the cylindrical elements are in serpentine patterns but out of phase with adjacent cylindrical elements. The particular pattern and how many undulations per unit of length around the circumference of the cylindrical element 12, or the amplitude of the undulations, are chosen to fill particular mechanical requirements for the stent such as radial stiffness.

The number of undulations may also be varied to accommodate placement of interconnecting elements 13, *e.g.* at the peaks of the undulations or along the sides of the undulations as shown in FIGS. 5 and 10.

The stent 10 of the present invention can be made

in many ways. However, the preferred method of making the stent is to coat a thin-walled tubular member, such as stainless steel hypotubing with a material which is resistive to chemical etchants and then to remove portions of the coating to expose underlying hypotubing which is to be removed but to leave coated portions of the hypotubing in the desired pattern for the stent so that subsequent etching will remove the exposed portions of the metallic tubing, but will leave relatively untouched the portions of the metallic tubing which are to form the stent. The coated portion of the metallic tube is in the desired shape for the stent. An etching process avoids the necessity of removing burrs or slag inherent in conventional or laser machine processes. It is preferred to remove the etchant-resistive material by means of a machine-controlled laser as illustrated schematically in FIG. 6.

A coating is applied to a length of tubing which, when cured is resistive to chemical etchants. "Blue Photoresist" made by the Shipley Company in San Jose, California, is an example of a suitable commercially available photolithographic coating. The coating is preferably applied by electrophoretic deposition.

To ensure that the surface finish is reasonably uniform, one of the electrodes used for the electrochemical polishing is a doughnut-shaped electrode which is placed about the central portion of the tubular member.

The tubing may be made of any suitable biocompatible material such as stainless steel, titanium, tantalum, superelastic NiTi alloys and even high strength thermoplastic polymers. The stent diameter is very small, so the tubing from which it is made must necessarily also have a small diameter. Typically the stent has an outer diameter on the order of about 1.5mm (0.06 inch) in the unexpanded condition, the same outer diameter of the hypotubing from which it is made, and can be expanded to an outer diameter of 2.5mm (0.1 inch) or more. The wall thickness of the hypotubing is about 0.076 mm (0.003 inch). In the instance when the stent is plastic, it has to be heated within the arterial site where the stent is expanded to facilitate the expansion of the stent. Once expanded, it is cooled to retain its expanded state. The stent may be conveniently heated by heating the fluid within the balloon or by heating the balloon directly by a suitable system such as disclosed in US-A-5,114,423 entitled "Dilation Catheter Assembly with Heated Balloon".

The stent may also be made of materials such as superelastic NiTi alloys such as described in US-A-5,411,476 entitled "Super-Elastic Guiding Member for Advancing Catheter".

In this case the stent would be formed full size but deformed (e.g. compressed) into a small diameter onto the balloon of the delivery catheter to facilitate transfer to a desired intraluminal site. The stress induced by the deformation transforms the stent from a martensite phase to an austenite phase and upon release of the force, when the stent reaches the desired intraluminal

location, allows the stent to expand due to the transformation back to the martensite phase.

Referring to FIG. 6, the coated tubing 21 is put in a rotatable collet fixture 22 of a machine controlled apparatus 23 for positioning the tubing 21 relative to a laser 24. According to machine-encoded instructions, the tubing 21 is rotated and moved longitudinally relative to the laser 24 which is also machine controlled. The laser selectively removes the etchant-resistive coating on the tubing by ablation and a pattern is formed such that the surface of the tube that is to be removed by a subsequent chemical etching process is exposed. The surface of the tube is therefore left coated in the discrete pattern of the finished stent.

A presently preferred system for removing the coating on the tubing includes the use of an 80 watt CO<sub>2</sub> laser, such as Coherent Model 44, in pulse mode (0.3 mS pulse length); 48 mA key current and 48 W key power with 0.75 W average power; at 100 Hz; Anorad FR=10; 1680 Pa; (12.5 Torr); with no assist gas. Low pressure air is directed through the fine focus head to ensure that no vapor contacts the lens. The assist gas jet assembly on the laser unit may be removed to allow a closer proximity of the fine focus head and the collet fixture. Optimum focus is set at the surface of the tubing. Cured photoresist coating readily absorbs the energy of the CO<sub>2</sub> wavelength, so that it can be readily removed by the laser. A coated 9.1 cm (4 inch) length of 1.5 mm (0.06 inch) stainless steel tubing is preferred and four stents can be patterned on the length of tubing. Three tabs or webs between stents provide good handling characteristics for the tubing after the etching process.

The process of patterning the resistive coating on the stent is automated except for loading and unloading the length of tubing. Referring again to FIG. 6 it may be done, for example, using a CNC opposing collet fixture 22 for axial rotation of the length of tubing, in conjunction with the CNC X/Y table 25 to move the length of tubing axially relative to a machine controlled laser as described. The entire space between collets can be patterned using the CO<sub>2</sub> laser set-up of the foregoing example. The program for control of the apparatus is dependent on the particular configuration used and the pattern to be ablated in the coating, but is otherwise conventional.

This process makes possible the application of present photolithography technology in manufacturing the stents. While there is presently no practical way to mask and expose a tubular photoresist-coated part of the small size required for making intravascular stents, the foregoing steps eliminate the need for conventional masking techniques.

After the coating is thus selectively ablated, the tubing is removed from the collet fixture 22. Next, wax such as ThermoCote N-4 is heated to preferably just above its melting point, and inserted into the tubing under vacuum or pressure. After the wax has solidified upon cooling, it is reheated below its melting point to allow

softening, and a small diameter stainless steel shaft is inserted into the softened wax to provide support. The tubing is then etched chemically in a conventional manner. After cutting the tabs connecting the stents any surface roughness or debris from the tabs is removed. The stents are preferably electrochemically polished in an acidic aqueous solution such as a solution of ELECTRO-GLO #300, sold by ELECTRO-GLO CO., Inc., in Chicago, IL, which is a mixture of sulfuric acid, carboxylic acids, phosphates, corrosion inhibitors and a biodegradable surface active agent. The bath temperature is maintained at about 43 to 57 degrees centigrade (about 110-135 degrees F.) and the current density is about 0.06 to 2.3 amps/cm<sup>2</sup> (about 0.4 to about 1.5 amps per in<sup>2</sup>). Cathode to anode area should be at least about two to one. The stents may be further treated if desired, for example by applying a biocompatible coating.

While the invention has been illustrated and described herein in terms of its use as an intravascular stent, it will be apparent to those skilled in the art that the stent can be used in other instances such as to expand prostatic urethras in cases of prostate hyperplasia. Other modifications and improvements can be made without departing from the scope of the invention.

#### Claims

1. A longitudinally flexible stent (10) formed from an elongated tubular member and comprising a plurality of cylindrically shaped elements (12) which are independently expandable in the radial direction and which are connected to one another by interconnecting elements (13) so as to be generally aligned on a common longitudinal axis such that, other than at an end of the stent (10), each cylindrically shaped element (12) has two adjacent cylindrically shaped elements (12) spaced in opposite axial directions, the or each interconnecting element (13) connecting one of said cylindrically shaped elements (12) to one of said adjacent cylindrically shaped elements (12) being circumferentially displaced from the or each interconnecting element (13) connecting said cylindrically shaped element (12) to the other of said adjacent cylindrically shaped elements (12).
2. The stent (10) of claim 1, wherein the cylindrically shaped elements (12) are adapted to retain their expanded condition upon the expansion thereof.
3. The stent (10) of claim 1 or claim 2, wherein the radially expandable cylindrically shaped elements (12), in an unexpanded condition, have an axial length which is less than their diameter.
4. The stent (10) of any preceding claim, wherein each radially expandable cylindrically shaped element (12) is formed of a structural member which

extends in an undulating pattern about the longitudinal axis of the stent (10) and which, in cross-section, has an aspect ratio of less than two to one.

5. The stent (10) of claim 4, wherein the structural member is formed of a biocompatible material selected from the group consisting of stainless steel, tungsten, tantalum, superelastic NiTi alloys and thermoplastic polymers.
6. The stent (10) of any preceding claim wherein the circumferential displacement of the interconnecting elements (13) is of equal magnitude.
7. The stent (10) of any preceding claim, wherein there are up to four interconnecting elements (13) disposed between adjacent radially expandable cylindrically shaped elements (12).
8. The stent (10) of any preceding claim, wherein the cylindrically shaped elements (12) are formed of structural members in an undulating pattern.
9. The stent (10) of claim 8, wherein the undulating patterns of the structural members of the stent (10) are in phase.
10. The stent (10) of claim 9, wherein the interconnecting elements (13) of the stent (10) are all connected to either the peaks or the valleys of the undulating structural members.
11. The stent (10) of claim 8, wherein the undulating pattern of the structural members of one cylindrical element (12) is out of phase with the undulating structural members of an adjacent cylindrical element (12).
12. The stent (10) of claim 1, wherein the stent (10) is coated with a biocompatible coating.
13. A kit comprising:
  - (a) an elongated stent delivery catheter (11) having proximal and distal extremities, and an expandable member (14) on the distal extremity; and
  - (b) a longitudinally flexible stent (10) formed from an elongated tubular member, the stent (10) being adapted to be slidably mounted on to the expandable member (14) of said catheter (11) and comprising a plurality of cylindrically shaped elements (12) which are independently expandable in the radial direction and which are connected to one another by interconnecting elements (13) so as to be generally aligned on a common longitudinal axis such that, other than at an end of the stent (10), each cylindri-



cally shaped element (12) has two adjacent cylindrically shaped elements (12) spaced in opposite axial directions, the or each interconnecting element (13) connecting one of said cylindrically shaped elements (12) to one of said adjacent cylindrically shaped elements (12) being circumferentially displaced from the or each interconnecting element (13) connecting said cylindrically shaped element (12) to the other of said adjacent cylindrically shaped elements (12).

#### Patentansprüche

1. Längsflexibler Stent (10), der aus einem langgestreckten rohrförmigen Element gebildet ist und eine Mehrzahl zylindrisch geformter Elemente (12) aufweist, die in radialer Richtung unabhängig aufweitbar sind und die miteinander durch Verbindungselemente (13) verbunden sind, so daß sie im wesentlichen auf einer gemeinsamen Längsachse derart ausgerichtet sind, daß, außer am Ende des Stents (10), jedes zylindrisch geformte Element (12) zwei benachbarte, in entgegengesetzte Axialrichtungen mit Abstand angeordnete zylindrisch geformte Elemente (12) aufweist, wobei das oder jedes Verbindungselement (13), das eines der zylindrisch geformten Elemente (12) mit einem der benachbarten zylindrisch geformten Elemente (12) verbindet, in Umfangsrichtung von dem oder jedem Verbindungselement (13), das das zylindrisch geformte Element (12) mit dem anderen der benachbarten zylindrisch geformten Elemente (12) verbindet, versetzt ist.
2. Stent (10) nach Anspruch 1, in dem die zylindrisch geformten Elemente (12) dazu ausgelegt sind, nach deren Aufweitung ihren aufgeweiteten Zustand zu behalten.
3. Stent (10) nach Anspruch 1 oder Anspruch 2, in dem die radial aufweitbaren zylindrisch geformten Elemente (12) im nicht aufgeweiteten Zustand eine axiale Länge aufweisen, die geringer als ihr Durchmesser ist.
4. Stent (10) nach einem der vorhergehenden Ansprüche, in dem jedes radial aufweitbare zylindrisch geformte Element (12) aus einem Strukturelement gebildet ist, das sich in einem Wellenmuster um die Längsachse des Stents (10) erstreckt, und das im Querschnitt ein Streckungsverhältnis von weniger als 2 : 1 aufweist.
5. Stent (10) nach Anspruch 4, in dem das Strukturelement aus einem biokompatiblen Material gebildet ist, gewählt aus der Gruppe, die aus rostfreiem Stahl, Wolfram, Tantal, superelastischen NiTi-

Legierungen und thermoplastischen Polymeren besteht.

6. Stents (10) nach einem der vorhergehenden Ansprüche, in dem der Umfangsversatz der Verbindungselemente (13) gleichmäßig groß ist.
7. Stent (10) nach einem der vorhergehenden Ansprüche, in dem bis zu vier Verbindungselemente (13) zwischen benachbarten radial aufweitbaren zylindrisch geformten Elementen (12) angeordnet sind.
8. Stent (10) nach einem der vorhergehenden Ansprüche, in dem die zylindrisch geformten Elemente (12) aus Strukturelementen in einem Wellenmuster gebildet sind.
9. Stent (10) nach Anspruch 8, in dem die Wellenmuster der Strukturelemente des Stents (10) in Phase liegen.
10. Stent (10) nach Anspruch 9, in dem die Verbindungselemente (13) des Stents (10) alle entweder mit den Bergen oder den Tälern der gewellten Strukturelemente verbunden sind.
11. Stent (10) nach Anspruch 8, in dem das Wellenmuster der Strukturelemente eines zylindrischen Elements (12) außer Phase mit den Wellenstrukturelementen eines benachbarten zylindrischen Elements (12) ist.
12. Stent (10) nach Anspruch 1, in dem der Stent (10) mit einer biokompatiblen Beschichtung beschichtet ist.
13. Ausrüstung, umfassend:
  - (a) einen langgestreckten stentausgabekatheter (11) mit einem proximalen und einem distalen Ende und einem aufweitbaren Element (14) an dem distalen Ende; und
  - (b) einen längsflexiblen Stent (10), der aus einem langgestreckten rohrförmigen Element gebildet ist, wobei der Stent (10) zum verschiebbaren Anbringen auf dem aufweitbaren Element (14) des Katheters (11) ausgelegt ist und eine Mehrzahl zylindrisch geformter Elemente (12) aufweist, die in radialer Richtung unabhängig aufweitbar sind und die miteinander durch Verbindungselemente (13) verbunden sind, so daß sie im wesentlichen auf einer gemeinsamen Längsachse derart ausgerichtet sind, daß, außer am Ende des Stents (10), jedes zylindrisch geformte Element (12) zwei benachbarte, in entgegengesetzte Axialrichtungen mit Abstand angeordnete zylindrisch

geformte Elemente (12) aufweist, wobei das oder jedes Verbindungselement (13), das eines der zylindrisch geformten Elemente (12) mit einem der benachbarten zylindrisch geformten Elemente (12) verbindet, in Umfangsrichtung von dem oder jedem Verbindungselement (13), das das zylindrisch geformte Element (12) mit dem anderen der benachbarten zylindrisch geformten Elemente (12) verbindet, versetzt ist.

#### Revendications

1. Sonde dilatable (10) dans la direction longitudinale formée d'un élément tubulaire allongé et comprenant une pluralité d'éléments de forme cylindrique (12) qui sont expansibles indépendamment dans la direction radiale et qui sont reliés entre eux par des éléments de liaison (13) afin d'être en général alignés le long d'un axe longitudinal commun, de manière que chaque élément de forme cylindrique (12) soit adjacent à deux éléments de forme cylindrique (12) espacés dans des directions axiales opposées, sauf à une extrémité de la sonde dilatable (10), le ou chaque élément de liaison (13), reliant un desdits éléments de forme cylindrique (12) à l'un desdits éléments de forme cylindrique adjacents (12), étant décalé le long de la circonférence par rapport à l'élément ou à chaque élément de liaison (13) reliant ledit élément de forme cylindrique (12) à l'autre desdits éléments de forme cylindrique adjacents (12).
2. Sonde dilatable (10) selon la revendication 1, dans lequel les éléments de forme cylindrique (12) sont aptes à garder leur état expansé après leur expansion.
3. Sonde dilatable (10) selon la revendication 1 ou la revendication 2, dans lequel les éléments de forme cylindrique (12) expansibles, dans la direction radiale, ont à l'état non-expansé une longueur axiale inférieure à leur diamètre.
4. Sonde dilatable (10) selon l'une quelconque des revendications précédentes, dans lequel chaque élément de forme cylindrique (12), expansible dans la direction radiale, est formé d'un élément structural qui s'étend en une configuration ondulante le long de l'axe longitudinal de la sonde dilatable (10) et qui a une section transversale dont le rapport des dimensions est inférieur à 2 : 1.
5. Sonde dilatable (10) selon la revendication 4, dans lequel l'élément structural est formé en une matière biocompatible choisie dans le groupe comprenant : l'acier inoxydable, le tungstène, le tantale, des alliages super élastiques de NiTi et des polymères ther-

moplastiques.

6. Sonde dilatable (10) selon l'une quelconque des revendications précédentes, dans laquelle le décalage le long de la circonférence des éléments de liaison (13) a une amplitude constante.
7. Sonde dilatable (10) selon l'une quelconque des revendications précédentes, dans laquelle il y a jusqu'à quatre éléments de liaison (13) placés entre des éléments de forme cylindrique (12) expansibles dans la direction radiale.
8. Sonde dilatable (10) selon l'une quelconque des revendications précédentes, dans laquelle les éléments de forme cylindrique (12) sont formés d'éléments structuraux de configuration ondulante.
9. Sonde dilatable (10) selon la revendication 8, dans laquelle les configurations ondulantes des éléments structuraux de la sonde dilatable (10) sont en phase.
10. Sonde dilatable (10) selon la revendication 9, dans laquelle les éléments de liaison (13) de la sonde dilatable (10) sont tous reliés soit aux pics soit aux creux des éléments structuraux ondulants.
11. Sonde dilatable (10) selon la revendication 8, dans laquelle la configuration ondulante des éléments structuraux d'un élément cylindrique (12) est déphasée par rapport aux éléments structuraux ondulants d'un élément cylindrique adjacent (12).
12. Sonde dilatable (10) selon la revendication 1, dans laquelle la sonde dilatable (10) est recouverte d'un enduit biocompatible.
13. Ensemble comprenant :
  - (a) un cathéter (11) de mise en place d'une sonde dilatable allongée ayant une extrémité proximale et une extrémité distale, ainsi qu'un élément expansible (14) à l'extrémité distale ; et
  - (b) une sonde dilatable (10) dans la direction longitudinale formée d'un élément tubulaire allongé, la sonde dilatable (10) étant apte à être montée coulissante sur l'élément expansible (14) dudit cathéter (11) et comprenant une pluralité d'éléments de forme cylindrique (12) qui sont expansibles indépendamment dans la direction radiale et qui sont reliés entre eux par des éléments de liaison (13) afin d'être en général alignés le long d'un axe longitudinal commun, de manière que chaque élément de forme cylindrique (12) soit adjacent à deux éléments de forme cylindrique (12) espacés dans

des directions axiales opposées, sauf à une  
extrémité de la sonde dilatable (10), le ou cha-  
que élément de liaison (13), reliant un desdits  
éléments de forme cylindrique (12) à l'un des-  
dits éléments de forme cylindrique adjacents 5  
(12), étant décalé le long de la circonférence  
par rapport à l'élément ou à chaque élément de  
liaison (13) reliant ledit élément de forme cylin-  
drique (12) à l'autre desdits éléments de forme  
cylindrique adjacents (12). 10

15

20

25

30

35

40

45

50

55

FIG. 1

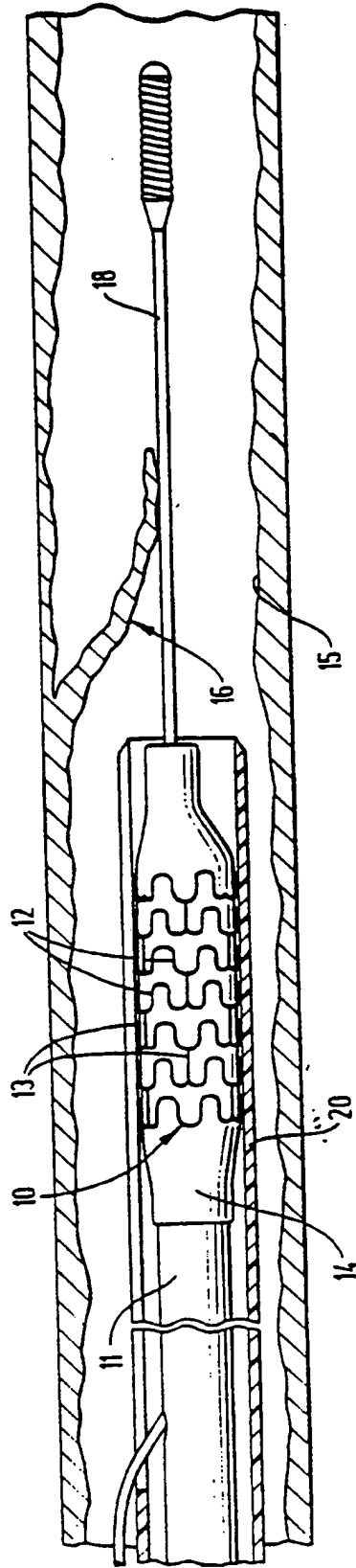
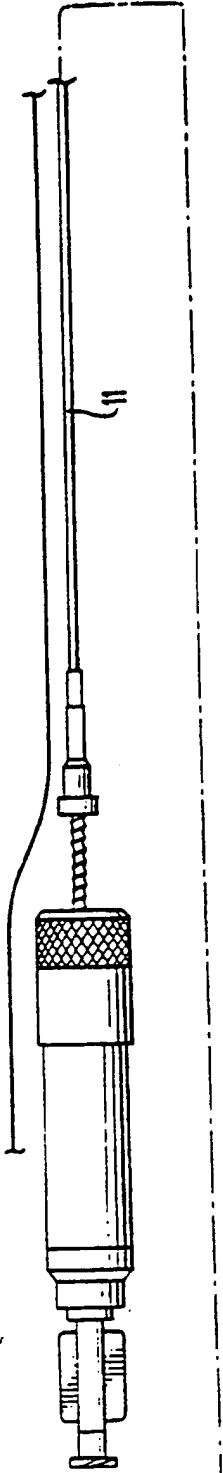


FIG. 3

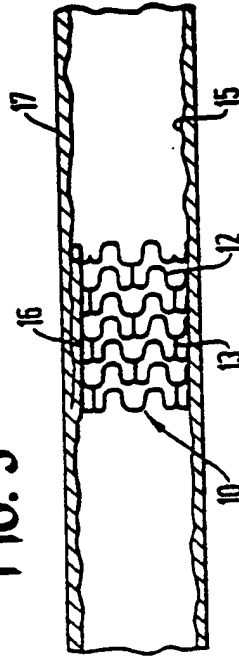
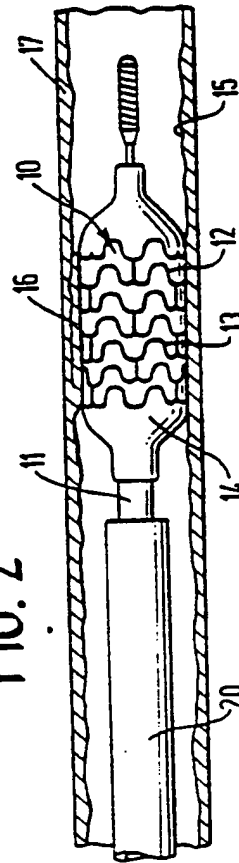


FIG. 2



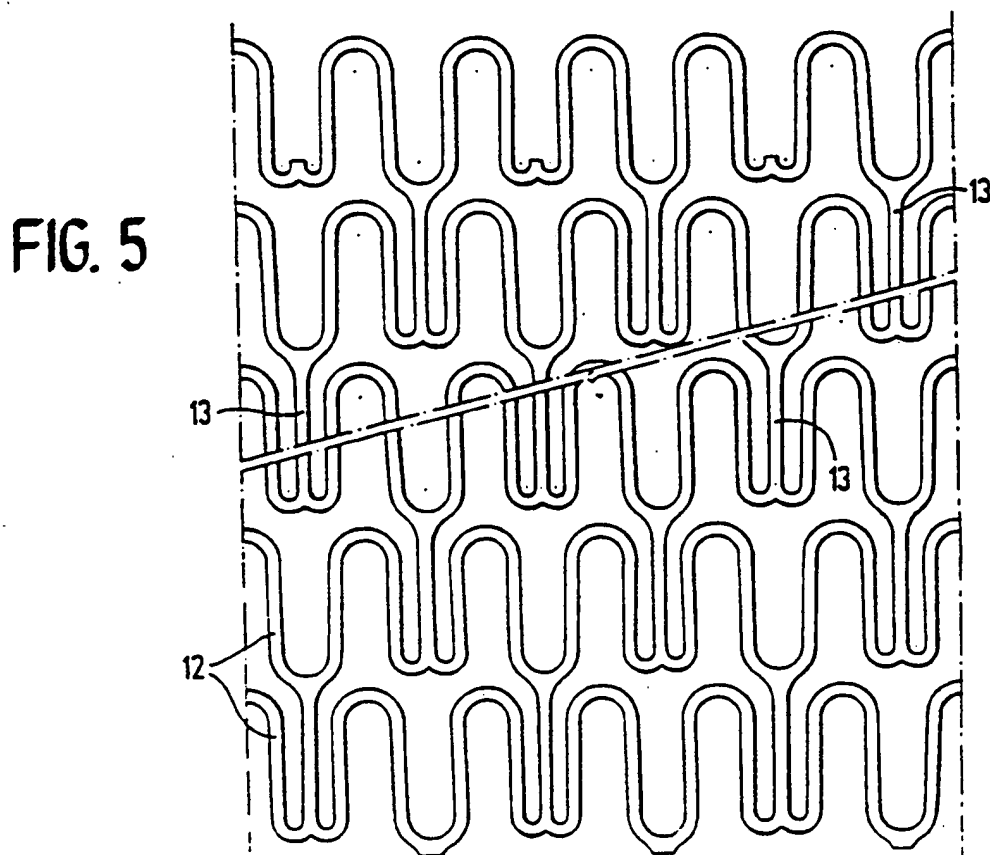
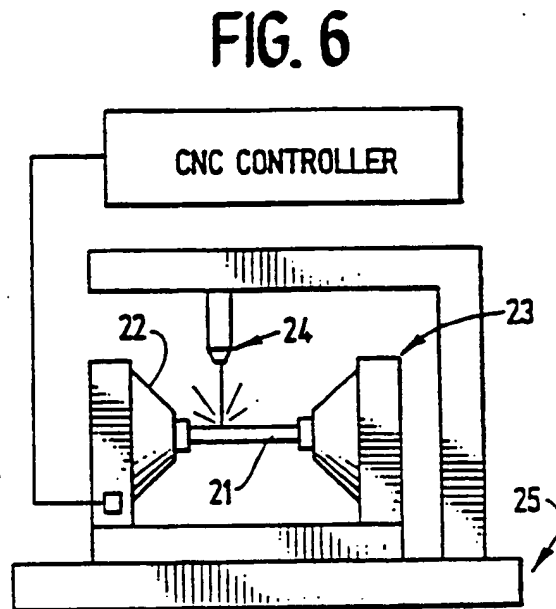
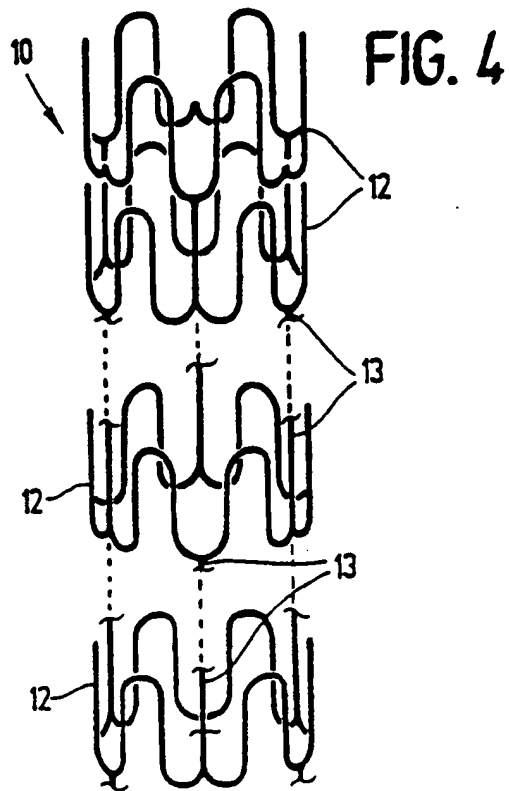


FIG. 7

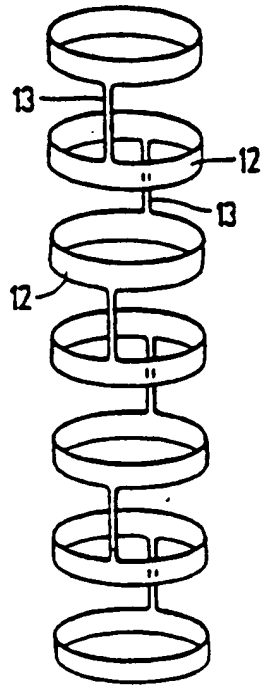


FIG. 8

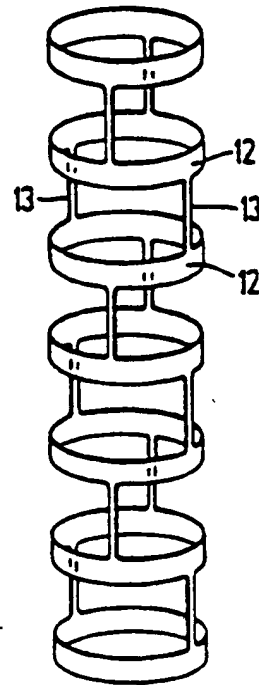


FIG. 9

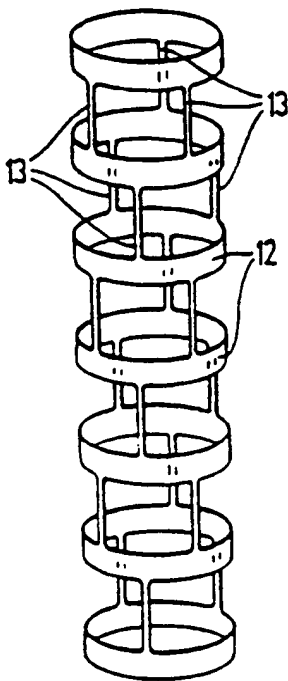


FIG. 10

